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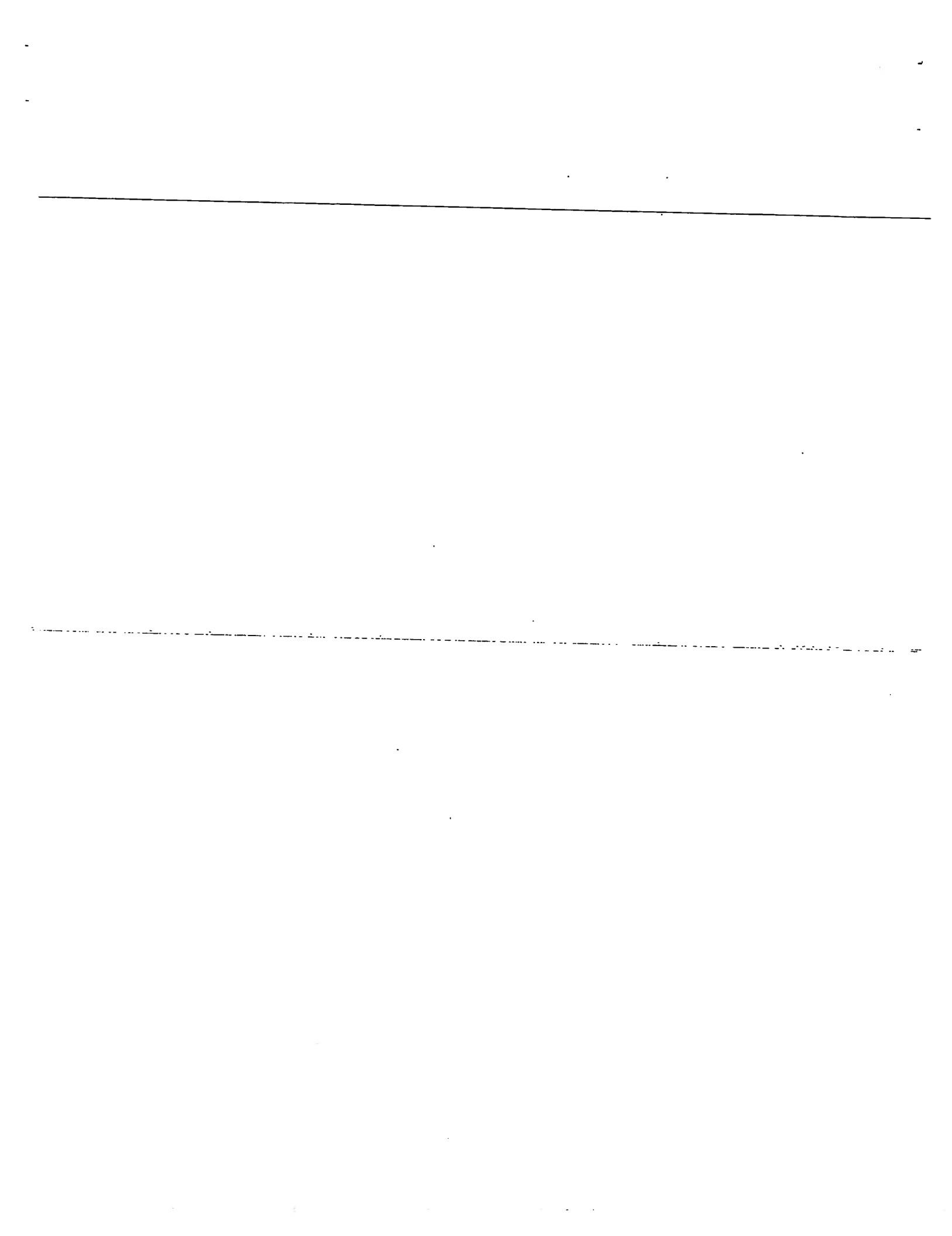
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Der Präsident des Europäischen Patentamts;
Im Auftrag

For the President of the European Patent Office
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Umicore
Rue du Marais 31
1000 Bruxelles
BELGIQUE

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Recovery of non-ferrous metals from zinc residues

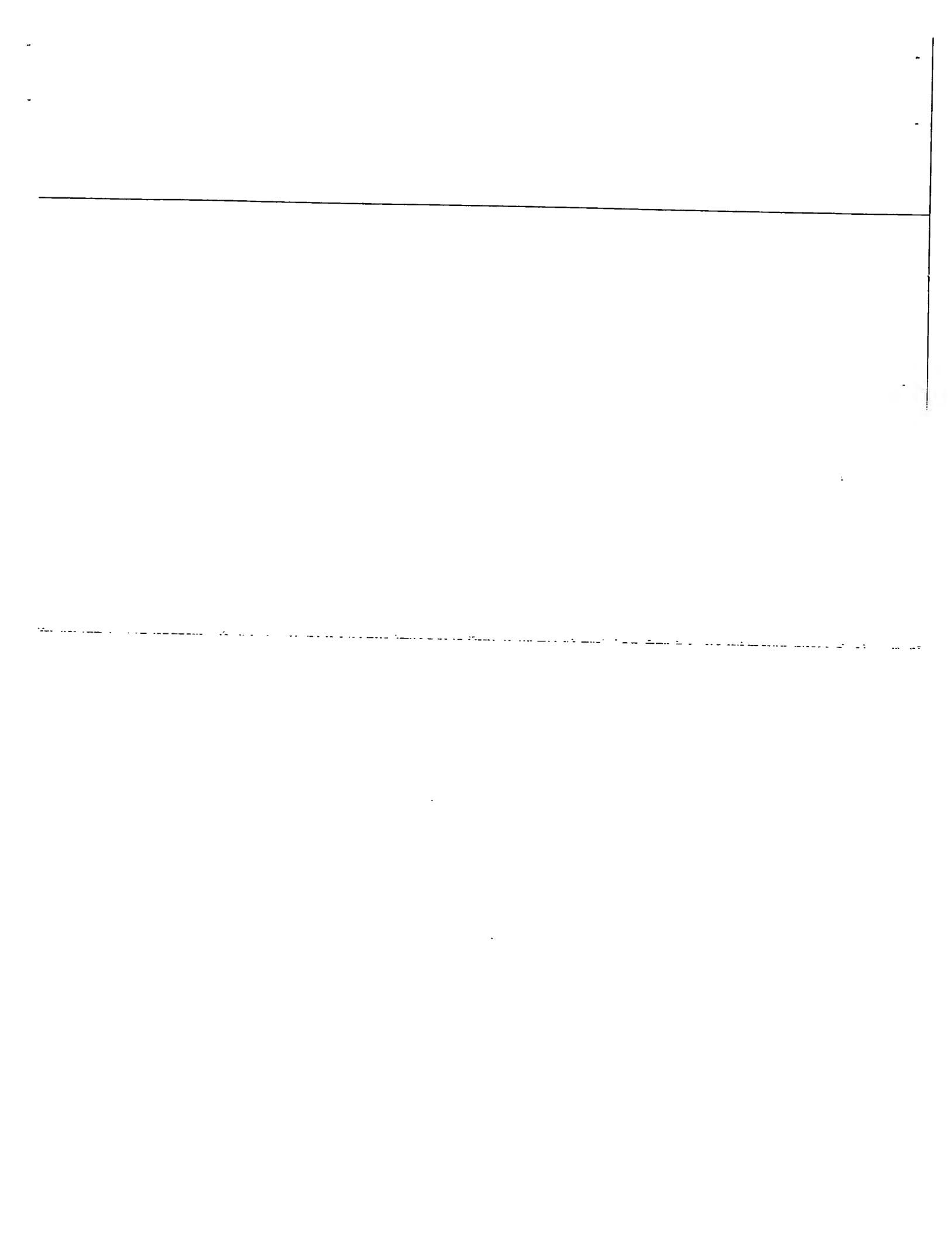
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Recovery of non-ferrous metals from zinc residues

5 This invention relates to the recovery of non-ferrous metals from zinc-bearing residues, in particular from residues produced by the zinc manufacturing industry.

Blende, which is an impure ZnS ore, is the main starting material for the production of Zn. The typical industrial practice encompasses an oxidative roasting step, producing ZnO together with sulphates or 10 oxides of the impurities. In subsequent steps, the ZnO in roasted blende is brought into solution by leaching in neutral conditions or in weak acidic conditions, thereby producing Zn-depleted residues, respectively referred to as neutral leach residue and as weak acid 15 leach residue in this description. However, during roasting, part of the Zn reacts with Fe, a typical impurity present in blende, and forms relatively insoluble zinc ferrite. The leach residues therefore contain, besides lead sulphate, calcium sulphate and other impurities, a sizeable fraction of Zn in the form of ferrite. According to present practice, the recovery of the Zn from ferrite 20 requires a specific hydro-metallurgical residue treatment using high acid concentrations of 50 to 200 g/l H₂SO₄. A disadvantage of this acidic treatment is that besides Zn, almost all the Fe and also other impurities such as As, Cu, Cd, Ni, Co, Tl, Sb are dissolved. As even low concentrations of these elements interfere with the subsequent 25 electrowinning of Zn, they must be removed from the zinc sulphate solution. While Cu, Cd, Co, Ni and Tl are precipitated by addition of Zn powder, Fe is typically discarded as hematite, jarosite or goethite through hydrolysis. Due to the danger of washout of heavy metals, these Fe-bearing residues have to be disposed off in a well-controlled landfill. Land-filling of such residues has however come 30 under heavy environmental pressure, rendering the sustainability of the process questionable. Another drawback of the above treatment is the loss of metals such as In, Ge, Ag and Zn in the Fe-bearing residue.

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An alternative treatment of the ferrite bearing residues is applied in some plants, using Waelz kilns, which produce a slag and a Zn and Pb containing fume. Similarly, a rotary flame-fired furnace of the Dorschel type can be used in a batch process. In still another 40 approach, the leach residue is processed, using coke as fuel, in a half shaft blast furnace, producing a Zn and Pb containing fume, matte and slag. These pyro-metallurgical treatments generally result

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in an excellent recovery of Zn and Pb, and, for some of them, in a significant recovery of Ag, Ge and In.

- These processes are however inadequate for modern zinc smelters, as 5 they cannot be scaled up to large single-vessel operations. By this fact, they are not a cost efficient solution for today's Zn smelters.

- Although numerous Zn fuming processes have been described, recent literature concentrates on the treatment of Zn-containing Fe-based 10 secondary residues, such as EAF dusts. In this respect, the production of DRI (Direct Reduced Iron) by the direct reduction process, allows for the reduction of metals in the solid phase, thus avoiding the extreme temperatures needed to produce a melt. Processing Fe-rich residues commonly aims at the recovery of pure 15 metallic Fe. This process is well suited for treating an Fe-rich feed such as EAF dust, but fails to produce a good quality DRI from the low-Fe, highly contaminated residues which are the subject of this invention. In such a case, the produced DRI indeed locks metals such as Cu, Ag and Ge and is therefore unsuitable for further processing 20 by the steel industry.

- The primary aim of the invention is to provide for a process for the separation and recovery of a wide range of non-ferrous metals like Cu, Ag, Ge and Zn from residues of the Zn manufacturing industry; 25 moreover, the process must ensure an environmentally acceptable output for Fe. The actual valorisation of Fe is however unimportant due to its relatively low concentration in the contemplated residues and to its rather low intrinsic value. These aims are achieved by a process for the valorisation of metal values in a Zn-, Fe- and Pb-bearing residue, comprising the steps of:
- subjecting the residue to a direct reduction step, thereby producing a reduced Fe-bearing phase and Zn- and Pb-bearing first fumes;
 - extracting the Zn- and Pb-bearing first fumes and valorising Zn and Pb;
 - subjecting the reduced Fe-bearing phase to an oxidising smelting step, thereby producing an Fe-bearing slag and second metals-bearing fumes;
 - extracting the second metals-bearing fumes and valorising at least 40 part of their metallic content.

The Zn-, Fe- and Pb-bearing residue may be a neutral leach residue or a weak acid leach residue. The fluidity of the slag can be enhanced

by avoiding massive formation of Fe_2O_3 , thus limiting the oxidation to the formation of mainly FeO . The presence of suitable amounts of an acidic flux such as sand and, preferably, of a basic flux such as lime, limestone or dolomite, also enhances the slag fluidity, thus

- 5 allowing for a lower working temperature.

If present, the major part of the Cu and Ag from the reduced Fe-bearing phase may be collected in a separate Cu-bearing phase during the oxidising smelting step.

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If the Zn-, Fe- and Pb-bearing residue also contains Ge, the fraction of the Ge present in said first fumes, produced in the direct reduction step, may be separated and forwarded to the oxidising smelting step. This separation of Ge is preferably performed by co-precipitation with Fe hydroxide or by addition of tannic acid. The same principles apply for In.

The first fumes may advantageously be oxidised in the reactor used for the direct reduction step.

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The reactor used for the direct reduction step can be a multiple hearth furnace; the reactor for the oxidising smelting step can be a submerged lance furnace.

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The details of the invention are now discussed. The first step consists of the reduction of the residue, after addition of reductants such as natural gas, coal or cokes, and possibly fluxes such as limestone ($CaCO_3$) to avoid sintering of the load, thereby producing Zn- and Pb-bearing fumes and a DRI-bearing phase

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containing, besides Fe, valuable metals such as Cu and Ag. Through leaching of these fumes, Zn and Pb can be separated in a Pb-containing residue and a Zn-containing leaching liquor. Ge, if present, is also dissolved in the leaching liquor, and can be precipitated as a Ge-containing residue and fed to the oxidising furnace, further enhancing the global Ge-recovery.

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The DRI-bearing phase is smelted and oxidised in a second furnace, preferably after addition of limestone and sand. The oxidation of Fe to FeO , and possibly to Fe_2O_3 , is particularly exothermic and brings in all the needed enthalpy. The process remains autogenous, even when e.g. 10% of the total feed consists of Ge-containing residues with moisture content of around 25%. However, if more than 50% of the Fe is oxidised to Fe_2O_3 instead of to FeO , the rise in liquidus

temperature of the slag entails disadvantages such as increased lining wear and higher energy demands. The fact that Fe is slagged instead of being recovered as a metal, is a small price to pay for achieving an excellent separation and recovery of the non-ferrous metals. The slag appears to be environmentally acceptable and can even be upgraded as gravel substitute in concrete. Next to this slag, two other streams can be recovered in this operation: fumes, typically containing Ge and In, and a Cu-bearing phase containing precious metals such as Ag. This phase can be refined in a classical Cu or precious metals flowsheet, separating the Cu and Ag in a Cu-rich and a Ag-rich stream. The fumes can be leached and used as feedstock for the production of Ge.

Useful reactor types such as multiple hearth furnaces and submerged lance furnaces lend themselves to the large single-vessel operations described in this invention. The overall process is energy efficient, bears low operational costs and ensures a high metal value recovery. The invention thus provides for an essentially waste-free process that can compete economically with hydro-metallurgical Zn residue treatments.

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The following example illustrates the separation of different non-ferrous metals contained in a roasted and subsequently leached blend. The analysis of the different feeds and productions are given in Table 1, together with the material balance.

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10.000 g of weak acid leach residue, which mainly consists of zinc ferrite ($ZnO \cdot Fe_2O_3$), lead sulphate ($PbSO_4$), calcium sulphate ($CaSO_4$), zinc sulphate ($ZnSO_4$) and impurities like CaO , SiO_2 , MgO , Al_2O_3 , Cu_2O , SnO , is dried and thoroughly mixed with 3.000 g cokes, having a purity of >85% C. The amount of cokes is calculated in order to reduce Fe and the other metals in the DRI, and fume Zn and the other metals in the fumes, with an excess of around 5% to handle fluctuations in the feed composition. The metal content of the leach residue is given in Table 1, where "Others" refers to the impurities summed up above and to the bound oxygen and sulphur. For the cokes, "Others" refers to its ash content. No flux was added in this experiment.

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The mixture is then loaded in a direct reduction furnace with an internal diameter of 454 mm and a height of 470 mm. Two arms are perpendicularly attached to a central shaft. The shaft is rotating at 2 rpm. Four teeth are located on each arm, which continuously stir the load. The teeth are positioned in such a way that one arm pushes

- the load towards the central shaft, while the other arm pushes in the opposite direction, resulting in a continuous stirring of the load without piling up at the furnace periphery. The atmosphere in the furnace is controlled by using a separate combustion chamber, where a mixture of air and natural gas is combusted. The nominal gas flow is 6 Nm³/hr natural gas and 47 Nm³/hr air, leading to a lambda value of 1. During 2 hours, the mixture is gradually heated until the temperature of the furnace atmosphere reaches 1100 °C; this temperature is maintained for approximately 4 hours. Fumes are collected using a cartridge filter, which is emptied every hour. The analysis of the fumes shown in Table 1 is the mean analysis of the hourly collected samples. These fumes are rich in Zn and Pb, present as ZnO and PbO/PbSO₄. "Others" in Table 1 mainly consists of oxygen and sulphur bound to the fumed metals, together with some carry-over.
- At the end of the reduction step, the DRI is removed from the furnace and sampled for analysis. The DRI obtained contains 88% of the Ag, 99% of the Cu, 70 % of the Ge, 7% of the Zn and 98% of the Fe when related to the feed. The low amount of Fe in the DRI (41,9%) is caused by the dilution with cokes remnants and light-metal oxides such as CaO, SiO₂, Al₂O₃, and MgO from the residue. This differentiates the obtained DRI from classic DRI's, which typically contain 80 to 90% Fe.
- In a second step, sand and limestone are added to the DRI and the resulting mixture is loaded in an oxidising furnace. It is melted under N₂-atmosphere in a crucible and maintained at 1200 °C for about 3 hours, while air is bubbled through the molten bath, resulting in the burning of the remaining C from the cokes and in the oxidation of Fe to FeO. The amount of air needed for oxidation of the Fe to FeO is calculated based on the analysis of the DRI given in Table 1. Production of Fe₂O₃ is avoided to obtain a slag with as low a liquidus temperature as possible. The burning of cokes and the conversion of Fe to FeO generate enough energy to operate the furnace without any O₂ enrichment.
- After the oxidative smelting step, three separate phases were recovered: fumes, slag and a Cu-bearing phase. The fumes are collected in a cartridge filter. Table 1 shows that these fumes are again rich in Zn and Pb. Thanks to the low total amount of fumes, elements like Ge and In, but also impurities like As, Sn, Bi and Sb, undergo a substantial concentration when related to the feed.

- The environmental stability of the slag was tested on the slag as such and after formation of concrete containing 30% slag and 10% cement. The tests were performed according to European norm NEN 7343, whereby the material is broken to a PSD of less than 4 mm and percolated with acidified water. The leachability was assessed according to the VLAREA ("Vlaams reglement voor afvalvoorkoming") norm for non-ferro metallurgical slags. The leachability of both the slag and the slag-containing concrete proved to be well below the limits applicable to products intended for the building industry.
- The Cu-bearing phase and the slag are separated by decantation, sampled and analysed. The slag analysis shows minimal amounts of leachable heavy metals, such as Pb, ensuring that an environmentally clean slag is obtained. The high percentage of "Others" in the slag is attributable to CaO, SiO₂, MgO and Al₂O₃.

The Cu-alloy contains 89% of the Ag and 80% of the Cu in the feed, together with some impurities, mainly Fe, As, Sb, Sn.

Table 1: Material balance and metal recovery per produced phase

Component	Mass (g)	Feed to reduction step								
		Composition (wt.%)								
		Ag	Pb	Cu	Zn	Fe	S	Ge	C	Others
Residue	10000	0,04	4,7	2,2	27,0	23,0	4,4	0,008		38,7
Cokes	3000					0,2	0,4		89	10,4

Component	Mass (g)	Products of reduction step								
		Composition (wt.%)								
		Ag	Pb	Cu	Zn	Fe	S	Ge	C	Others
Pure DRI	5440	0,07	0,64	4,0	3,8	41,9	2,9	0,011	10	36,7
First fumes	4670	0,01	9,3	0,04	53,4	0,60	6,3	0,005		30,3

Component	Mass (g)	Feed to oxidising smelting step								
		Composition (wt.%)								
		Ag	Pb	Cu	Zn	Fe	S	Ge	C	Others
Pure DRI	5440	0,07	0,64	4,0	3,8	41,9	2,9	0,011	10	36,7
Sand	1220									100
Limestone	800									100

Component	Mass (g)	Products of oxidising smelting step								
		Composition (wt.%)								
		Ag	Pb	Cu	Zn	Fe	S	Ge	C	Others
Fe-slag	6780	<0,01	0,10	0,6	2,2	33,4	1,5	<0,001	<0,05	62,2
Cu-alloy	207	1,8	<0,1	85	<0,1	5,0	2,0	<0,001	<0,05	5,23
Second fumes	186	<0,01	15,2	1,0	32,4	1,0	2,35	0,32		47,7

(Table 1 continued)

Component	Fraction (wt. %)	Recovery (%)						
		Ag	Pb	Cu	Zn	Fe	S	Ge
Fe-slag	52	"	1	18	5	98	23	
Cu-alloy	2	89		80			1	
First fumes	36	11	93	1	92	1	65	30
Second fumes	1		6	1	2		11	70

- 5 The invented process thus achieves the separation of the metals as follows:
- Zn, Pb and a minor part of the Ge in first fumes which can be treated by known means for separation of Pb and Ge in different residues, and of Zn in a leach liquor;
- 10 10 - Cu and Ag in an alloy which can be refined using a classical Cu and precious metals flowsheet;
- Ge in second fumes that are greatly Ge-enriched and from which Ge can be refined cost-effectively;
 - Fe in an inert, environmentally clean slag, reusable as e.g. gravel substitute in concrete.
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CLAIMS

1. Process for the valorisation of metal values in a Zn-, Fe- and Pb-bearing residue, comprising the steps of:
 - subjecting the residue to a direct reduction step, thereby producing a reduced Fe-bearing phase and Zn- and Pb-bearing first fumes;
 - extracting the Zn- and Pb-bearing first fumes and valorising Zn and Pb;
 - subjecting the reduced Fe-bearing phase to an oxidising smelting step, thereby producing an Fe-bearing slag and second metals-bearing fumes;
 - extracting the second metals-bearing fumes and valorising at least part of their metallic content.
2. Process according to claim 1, wherein the Zn-, Fe- and Pb-bearing residue is a neutral leach residue or a weak acid leach residue.
3. Process according to claims 1 or 2, characterised in that during the oxidising step, Fe is oxidised to mainly FeO in the slag.
4. Process according to claim 3, characterised in that an acidic flux and, preferably, also a basic flux are present in the oxidising smelting step.
5. Process according to claims 1 or 4, characterised in that the Zn-, Fe- and Pb-bearing residue contains Cu and Ag, and that, during the oxidising smelting step, a separate Cu-alloy phase is produced containing a major part of the Cu and Ag.
6. Process according to any one of claims 1 to 5, characterised in that the Zn-, Fe- and Pb-bearing residue contains Ge, and that, after the direct reduction step, the fraction of the Ge present in the first fumes is separated and forwarded to the oxidising smelting step.
7. Process according to claim 6, whereby the separation of Ge is performed by co-precipitation with Fe hydroxide or by addition of tannic acid.
8. Process according to any one of claims 1 to 7, whereby the first fumes are oxidised in the reactor used for the direct reduction step.

9. Process according to any one of claims 1 to 8, whereby the reactor used for the direct reduction step is a multiple hearth furnace.
- 5 10. Process according to any one of claims 1 to 9, whereby the reactor used for the oxidising smelting step is a submerged lance furnace.

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ABSTRACTRecovery of non-ferrous metals from zinc residues

- 5 The invention relates to a process for the separation and recovery of non-ferrous metals from zinc-bearing residues, in particular from residues produced by the zinc manufacturing industry.
- 10 The process allows for the valorisation of metal values in a Zn-, Fe- and Pb-bearing residue, and comprises the steps of:
- subjecting the residue to a direct reduction step, thereby producing a reduced Fe-bearing phase and Zn- and Pb-bearing first fumes;
 - extracting the Zn- and Pb-bearing first fumes and valorising Zn and Pb;
 - subjecting the reduced Fe-bearing phase to an oxidising smelting step, thereby producing an Fe-bearing slag and second metals-bearing fumes;
 - extracting the second metals-bearing fumes and valorising at least part of their metallic content.

The main advantage of this process is that an environmentally acceptable output for Fe is obtained.

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